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9	117	(436/827).CCLS.	USPAT;	2002/07/29 08:25
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12	14	sialo adj ganglioside	USPAT;	2002/07/29 08:27
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			EPO;	
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			DERWENT	
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			EPO;	
1			DERWENT	
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			EPO;	
			DERWENT	
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Search History 7/29/02 9:11:11 AM Page 1

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}			EPO;	
			DERWENT	
19	30	sialoadhesin	USPAT;	2002/07/29 08:31
			US-PGPUB;	
			EPO;	
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		neuac or sialoadhesin or siglec	EPO;	
		Heads of Stated affecting of Styles	DERWENT	
22	658097	complex\$2	USPAT;	2002/07/29 08:46
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			EPO;	
			DERWENT	
23	12	(sialoganglioside or (sialo adj ganglioside) or sialolipid or	USPAT;	2002/07/29 08:48
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			DERWENT	
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1		(sialo adj lipid) or (sialo adj tn) or (sialyl adj tn) or stn or	US-PGPUB;	
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00	4500	 	DERWENT	0000/07/00 00 07
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			EPO; DERWENT	
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30	13126	(multiple adj sclerosis) same (multiple adj sclerosis)	USPAT;	2002/07/29 09:08
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	[EPO;	
L	<u></u>		DERWENT	

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		neuac or sialoadhesin or siglec	EPO;	
		Trouble of blandarioum of bigliob	DERWENT	
3	658097	complex\$2	USPAT;	2002/07/29 09:41
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			EPO;	
			DERWENT	
4	12	(sialoganglioside or (sialo adj ganglioside) or sialolipid or	USPAT;	2002/07/29 09:41
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		or sialolipid or (sialo adj lipid) or (sialo adj tn) or (sialyl adj tn)	US-PGPUB;	2002/01/20 00:10
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			DERWENT	
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			EPO;	
1			DERWENT	
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			DERWENT	
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			DERWENT	
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'			EPO;	
			DERWENT	
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		stn or neuac or sialoadhesin or siglec)	EPO;	
	-	(400,040,0010, 1411	DERWENT	
14	3	((436/813).CCLS.) and (sialocomplex or (sialoganglioside or	USPAT;	2002/07/29 10:23
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L*** (129)SEA FILE=BIOSIS ABB=ON PLU=ON L*** AND L***

L*** (46)SEA FILE=CA ABB=ON PLU=ON L*** AND L***

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           153 L1 NOT 2002/PY
=> s 12 not 2001/py
           147 L2 NOT 2001/PY
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L<sub>3</sub>
AN
    1993:200903 BIOSIS
DN
     PREV199344097153
TI
     Seric immune complexes in multiple
     sclerosis do not contain MBP epitopes.
ΑU
    Geffard, Michel (1); Boullerne, Anne; Brochet, Bruno
CS
     (1) Lab. Immunologie Pathologie, Univ. Bordeaux II, BP 66, 146 rue Leo
     Saignat, 33076 Bordeaux Cedex France
    Brain Research Bulletin, (1993) Vol. 30, No. 3-4, pp. 365-368.
SO
    Meeting Info.: Satellite Symposium of the Third IBRO (International Brain
     Research Organization) World Congress of Neuroscience on Development,
     Plasticity, and Regeneration in the Spinal Cord: Cellular and Molecular
     Interactions Quebec, Quebec, Canada August 11-14, 1991
     ISSN: 0361-9230.
    Article
DT
LA
     English
=> d 13 21 bib ab
L3
    ANSWER 21 OF 147 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
AN
     1992:160579 BIOSIS
DN
     BR42:76779
ΤI
     ANTIBRAIN ANTIBODIES IN MULTIPLE SCLEROSIS PATIENTS.
    HENNEBERG A E; MAYLE D M; PFAU J; ELSNER U; KORNHUBER H H
ΑU
CS
    UNIV. ULM DEP. NEUROL., D-7900 ULM.
SO
     THIRD INTERNATIONAL CONGRESS ON NEUROIMMUNOLOGY, JERUSALEM, ISRAEL,
     OCTOBER 27-NOVEMBER 1, 1991. J NEUROIMMUNOL. (1991) 0 (SUPPL 1), 183.
     CODEN: JNRIDW. ISSN: 0165-5728.
DT
     Conference
FS
     BR; OLD
LΑ
     English
=> d 13 41 bib ab
    ANSWER 41 OF 147 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
L3
ΑN
     1987:189368 BIOSIS
DN
    BA83:97492
TΤ
     THE DETECTION OF BRAIN ANTIGENS WITHIN THE CIRCULATING IMMUNE
     COMPLEXES OF PATIENTS WITH MULTIPLE SCLEROSIS.
ΑU
     FRIEDMAN J; BUSKIRK D; MARINO L J JR; ZABRISKIE J B
CS
    DEP. BACTERIOL. IMMUNOL., ROCKEFELLER UNIV., NEW YORK, N.Y. 10021, USA.
SO
     J NEUROIMMUNOL, (1987) 14 (1), 1-18.
     CODEN: JNRIDW. ISSN: 0165-5728.
    BA: OLD
FS
LА
     English
AΒ
     Immune complexes isolated from sera of patients with
     multiple sclerosis (MS) were analyzed for their
     antigenic content. Immune complexes precipitated with
     polyethylene glycol were inoculated into rabbits. The antisera raised were
```

shown to react to MS and normal brain antigens by crossed immunoelectrophoresis, ELISA and nitrocellulose transfer. Additionally, these antisera reacted by co-precipitation with the measles nucleocapsid antigen. As persistent viral infection has not been shown to occur in MS, these studies support the possibility that brain antigens within the immune complex share antigenic determinants with a specific viral antigen seen earlier in life by the host.

=> d 13 43 bib ab

- L3 ANSWER 43 OF 147 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- AN 1986:399731 BIOSIS
- DN BA82:85211
- TI ANTI-GLYCOLIPID ANTIBODIES AND THEIR IMMUNE COMPLEXES IN MULTIPLE SCLEROSIS.
- AU KASAI N; PACHNER A R; YU R K
- CS DEP. NEUROLOGY, YALE UNIV. SCH. MED, 333 CEDAR ST., NEW HAVEN, CT 06510.
- SO J NEUROL SCI, (1986) 75 (1), 33-42. CODEN: JNSCAG. ISSN: 0022-510X.
- FS BA; OLD
- LA English
- Antibody titers against myelin constituents in sera and CSF of patients with multiple sclerosis (MS) were examined by a solid-phase radioimmunoassay. Anti-GM4 and anti-galactocerebroside antibody titers were significantly elevated in the CSF of MS patients, but not anti-GM1 and anti-myelin basic protein antibodies. In sera of MS patients, the titers of antibodies against these myelin constituents were not elevated. Total IgG level was also significantly elevated in the CSF, but not in the sera of MS patients. Immune complexes from the CSF of MS patients were dissociated by acid-ultrafiltration. These data suggest that antibodies of the IgG class against GM4 and galactocerebroside are present in CSF of MS patients, and a significant number of them exist as immune complexes with their corresponding glycolipid antigens.
- => d 14 bib ab
- L4 ANSWER 1 OF 1 CA COPYRIGHT 2002 ACS
- AN 112:18592 CA
- TI Anti-ganglioside GD1a monoclonal antibody MZ, MZ-producing cells, and MZ-containing reagent for diagnosis of cancer and systemic lupus erythematosus
- IN Shimada, Shizuo; Iwata, Daiji; Sato, Wakao
- PA Mitsui Toatsu Chemicals, Inc., Japan
- SO Eur. Pat. Appl., 13 pp.
 - CODEN: EPXXDW
- DT Patent
- LA English
- FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
		-			
ΡI	EP 307186	A2	19890315	EP 1988-308273	19880907
	EP 307186	A 3	19900314		
	EP 307186	B1	19940622		
	R: DE, FR,	GB			
	JP 01067198	A2	19890313	JP 1987-221862	19870907
	JP 07116238	B4	19951213		
	CA 1314246	A1	19930309	CA 1988-576560	19880906

	US	5192662	A	19930309	US	1988-241291	19880907
	JР	08187081	A2	19960723	JP	1995-167874	19950612
	JΡ	2635946	B2	19970730			
PRAI	JΡ	1987-221862		19870907			

ΑB Disclosed are a novel anti-ganglioside GD1a monoclonal antibody (MZ) which is capable of recognizing ganglioside GDla but is practically incapable of recognizing glycolipids GalCer, LacCer, Gb3, Gb4, GA1, GA2, GM1, GM2, GM3, GD1b, GT1b, GQ1b, Fuc-GM1, nLC4, and sialosyl nLC4; MZ-producing cells; an MZ-contg. reagent; and a method for the detection or quantification of GD1a using the reagent, e.g. to diagnose cancer, systematic lupus erythematosus (SLE), and diseases resulting from org. injury to the nervous system. Mice were immunized 4-15 times with formalin-treated Salmonella minnesota, and monoclonal antibodies were produced by std. techniques by fusion of their spleen cells with mouse myeloma NS-1 and screening of the clones. Serum from cancer or SLE patients was adsorbed on protein A-polystyrene, immune complexes were dissocd., the supernatant was extd. with CHCl3:MeOH (2:1), and the GD1a-contg. CHCl3 layer was dried and dissolved in buffered saline contg. 1% bovine serum albumin. This soln. was placed in microtiter plate wells and incubated with polystyrene-immobilized MZ for 3 h at 37.degree., biotinylated MZ for 1 h at room temp., and peroxidase-labeled avidin for 1 h at room temp. After treatment with substrate, the absorbance was read and compared to a std. curve for GD1a detn. Cancer and SLE patients had higher GD1a levels than healthy subjects.